# EXHIBIT 1

(19)日本国特許庁 (JP)

# (12) 公開特許公報(A)

(11)特許出願公開番号 特別平9-8178

(43)公開日 平成9年(1997)1月10日

(51) Int.Cl.6		識別記号	庁内整理番号	FΙ			技術表示箇所
H01L	23/29			HOIL	23/30	R	
	23/31			C 0 8 G	59/18	NJW	
C 0 8 G	59/18	NJW			59/26	NHQ	
	59/26	NHQ		C08L	63/00	NLC	
COSL	63/00	NLC					
	-			of the or	-D1:04:-D	TO SHOPEN	(A 0 W

		審查請求	未請求 請求項の数3 OL (全 6 頁)					
(21)出願番号	特顯平7-147661	(71)出題人	000005832					
(22)出顧日	松下電工株式会社 平成7年(1995) 6月14日 大阪府門真市大字門真1048番							
		(72)発明者	原 竜三 大阪府門真市大字門真1048番地松下電工株					
		(70) 70 77	式会社内					
		(72)発明者	官谷 至祥 大阪府門真市大字門真1048番地松下電工株 式会社内					
		(72)発明者	市川 貴之 大阪府門真市大字門真1048番地松下電工株 式会社内					
		(74)代理人	弁理士 佐藤 成示 (外1名) 最終頁に続く					

(54) 【発明の名称】 封止用エポキシ樹脂組成物及びそれを用いた半導体装置

# (57)【要約】

【目的】 耐湿性、耐吸湿半田クラック性、密着性に優れ、ポストキュアー後の変色を低減した緑色の封止用エポキシ樹脂組成物及びそれを用いた半導体装置を提供する

【構成】 1分子中に2個以上のエポキシ基を有するエポキシ樹脂に、硬化剤として1分子中に2個以上のフェノール性水板基を有するフェノール機能、無機充填剤、硬化促進剤及び額料を添加してなる封止用エポキシ樹脂組成物において、前記エポキシ樹脂として、ピフェニル型エポキシ樹脂を含有し、かつ、前記額料として、塩素化銅フタロシアニングリーンを封止用エポキシ樹脂組成か全をは対して0.05~1電量を及びカーボンブラックを封止用エポキシ樹脂組成物全量に対して0.07~02 重量%含有する。前記封止用エポキシ樹脂組成物の硬化物により封止してなる。

### 【特許請求の範囲】

【請求項1】 1分子中に2個以上のエポキシ基を有す るエポキシ樹脂に、硬化剤として1分子中に2個以上の フェノール性水酸基を有するフェノール樹脂、無機充填 剤 砂化促進剤及び顔料を添加してなる封止用エポキシ 樹脂組成物において、前記エポキシ樹脂として、下記の 一般式①で表されるビフェニル型エポキシ樹脂及び/▽ は下記の一般式②で表されるジシクロペンタジエン系エ\* \* ポキシ樹脂を含有し、かつ、前記願料として、塩素化銅 フタロシアニングリーンを封止用エポキシ樹脂組成物全 骨に対して0.05~1重量%及びカーボンブラックを 封止用エポキシ樹脂組成物全量に対して0.01~0. 2 重畳%含有することを特徴とする封止用エポキシ樹脂 組成物。

[(1:1]

(式中R,-R,は、H、CH,又はC,H,である。)

[化2]

#### (式中かは繰り返し単位を示す0~5の他)

【請求項2】 前記フェノール樹脂として、下記の一般 ※する請求項1記載の封止用エポキシ樹脂組成物。 式式②で表されるフェノール樹脂を含有することを特徴と※ [(1:3]

【請求項3】 請求項1又は請求項2記載の封止用エポ キシ樹脂組成物の硬化物により封止してなることを特徴 とする半遺体装置。

#### 【発明の詳細な説明】

### [0001]

【産業上の利用分野】本発明は、雷気部品、雷子部品、 半導体素子等を封止するための封止用エポキシ樹脂組成 40 物及びそれを用いた半導体装置に関する。

#### [0002]

【従来の技術】従来、ダイオード、トランジスター、集 精同路等の電気・電子部品や半導体装置等の封止方法と して、例えば、エポキシ樹脂やシリコン樹脂等による樹 脂封止方法や、ガラス、金属、セラミックス等を用いた ハーメチックシール法が採用されてきているが、近年で は、信頼性の向上とともに大量生産やコストの面でメリ ットのあるエポキシ樹脂を用いる封止法においては、ク

ノールノボラック型樹脂を硬化剤成分とする組成物から なる成形材料が最も一般的に使用されている。

【0003】 しかしながら、 1C、 LSI、 VLSI等 の電子部品や半導体装置の高密度化、高集積化にともな って、モールド樹脂の薄肉化が進んでおり、これまでの エポキシ樹脂組成物では、必ずしも満足に対応すること ができなくなっている。例えば、表面実装用デバイスに おいては、実装時にデバイス自身が半田に直接浸漬され る等、急激に高温苛酷環境下に曝されるため、パッケー ジクラックの発生が避けられない事情となっている。す なわち、成形後の保管中に吸湿した水分が、高温にさら される際に急激に気化膨張し封止樹脂がこれに耐えきれ ずに半導体装置のパッケージにクラックが生じる。ま た、一部のダイオードやトランジスターには、緑色が使 用されているものもあるが、封止用エポキシ樹脂組成物 は、一般的に黒色である。緑色は、イメージが良く、月 レゾールノボラック型樹脂を樹脂成分とし、かつ、フェ 50 立つので、封止用エポキシ樹脂組成物として使用したい が、通常の緑色の顔料を用いると、耐湿性が悪く、ポストキュアー後に変色するというような問題点があった。 【0004】

【発明が解決しようとする課題】本発明は前記の事情に 鑑みてなされたもので、その目的とするところは、耐湿 性、耐吸湿半田クラック性、密着性に優れ、ポストキュ ア一後の変色を低減した緑色の封止用エポキシ樹脂組成 物及びそれを用いた半導体装置を提供することにある。 【〇〇〇5】

【課題を解決するための手段】本発明の請求項1に係る 10 封止用エポキシ樹脂組成物は、1分子中に2個以上のエ ポキシ基を有するエポキシ樹脂に、硬化剤として1分子\*

\*中に2個以上のフェノール性水酸基を有するフェノール 樹脂、無機充填剤、硬化促進剤及び筋料を添加してなる 封止用エボキや樹脂組成物において、前起エボキや樹脂 として、下配の一般式ので表されるビフェニル型エボキ シ樹脂及び/又は下配の一般式ので表されるビフェニル型ンロベ ンタジエン系エボキや樹脂を含有し、かつ、前記顔料と して、塩素化銅ブタロシアニングリーンを封止用エボキ シ樹脂組成物全量に対して0.05~1重量%及びカー ボンブラックを封止用エボキシ樹脂組成物全量に対して 0.01~0.2重量%含有することを特徴とする。

[0006] [化4]

(式中R、-R。は、H、CH、又はGH、である。)

[0007]

#### (式中mは繰り返し単位を示す0~5の数)

【0008】本発明の請求項2に係る封止用エポキシ樹脂組成物は、前記フェノール樹脂として、下記の一般式 ②で表されるフェノール樹脂を含有することを特徴とす★

[0009]

式中R<sub>6</sub>-R<sub>6</sub>は、H、フェノール性水酸基、 ハロゲン原子又は炭素数1~5のアルキル基で、 nは頼り返し単位を示す1~5の数である。

【0010】本発明の請求項3に係る半導体装置、請求項1又は請求項2記載の封止用エポキシ樹脂組成物の硬 40 化物により封止してなることを特徴とする。

[0011] 以下、本発明を辞述する。本発明に用いる エポキシ樹脂は、1分子中に少なくとも2個のエポキシ 基を有するエポキシ樹脂であり、このエポキシ樹脂として、前記の一般式ので表されるビフェニル型エポキシ樹 脂及び/又は前記の一般式ので表されるジシクロペンタ ジエン系エポキシ樹脂を含有することが必須である。ま なわち、ピフェニル型エポキシ樹脂は剛直なピフェニル 骨格を有するため、得られる対止の低弾性率であっ 、高い熱時強度を持ち、密着力に優れたものになり、 ジシクロベンタジエン系エボキン樹脂を用いることにより、硬化物の吸湿性が小さくなる。つまり、前記エボキシ樹脂を用いることにより、吸煙率を下げ、リードフレームとの密着力を向上させることができ、耐湿性、耐半田クラック性に優れた硬化物が得られる封止用エボキシ樹類組被前になる。

【0012】本発明に用いる硬化剤としては、1分子中 にフェノール性水酸基を少なくとも2個有する硬化剤で あれば、いずれの硬化剤でも用いることができ、例え は、フェノールノボラック樹脂、クレゾールノボラック 樹脂及び多官郎フェノール樹脂等がある。

50 【0013】本発明では、無機充填材として、溶融シリ

カ、結晶シリカ、アルミナ及び発化ケイ素等を用いることができ、硬化促進剤として、ジアザビシクロウンデセン、トリフェールホスカニウム、テトラフェニルホスホニウム、テトラフェニルボレート、イミダゾール及び3数アミン等を用いることができる。また、必要に応じてエポキシシラン等のカップリング剤、プロ人にポキシ時間 脳及び三酸化アンチモン等の難燃剤、シリコーン可撓剤、カルナパワックス並びにステアリン酸等の離型剤を用いることができる。

【0015】本発明に係る封止用エボキシ樹脂組成物は、硬化剤であるフェノール樹脂として、前配の一般式ので表されるフェノール樹脂と含すすることが好ましい。すなわち、前配の一般式ので表されるフェノール樹脂を用いることにより、硬化物の吸湿性が小さくなり、密着力が向上し、耐湿性、耐半田クラック性に優れる封30止用エボキシ樹脂組成物が得られる。

【0017】以上のように、本発明によると、耐湿性、 耐吸湿半田クラック性、密着性に優れ、ポストキュアー 後の変色を低減した緑色の封止用エポキシ樹脂組成物及 40 びそれを用いた半導体装置が得られる。

#### [0018]

【作用】本発明の請求項1に係る封止用エポキシ樹脂組成物は、エポキシ樹脂として、前配の一般式ので表されるビフェニル型エポキシ樹脂及び/又は前配の一般式ので表されるジシクロベンタジエン系エポキシ樹脂を含有するので、ビフェニル型エポキシ樹脂が剛直なビフェニル骨格を有するため、ビフェニル型エポキシ樹脂を用いることにより、得ちれる封止品が原理性率であって、高い粉糖油酸で表れた。毎年かれたのにかり、ジン粉糖油酸を対ち、密やかに紹介されたのにかり、ジン粉糖油酸を対ち、密やかに紹介されたのにかり、ジン

ロベンタジエン系エボキン樹脂を用いることにより、硬化物の吸湿性が小さくなる。かつ、前記顔料として、塩素化銅フタロシアニングリーンを封止用エボキシ樹脂組成物全量に対して0.05〜1重銀%及びカーボンブラックを封止用エボキシ樹脂組成物全間に対して0.01〜0.2重量%含有するので、ボストキュアー後の変色を低減した緑色の封止用エボキシ樹脂組成物が得られ

[0019] 本発明の請求項2に係る封止用エポキシ樹 脂組成物は、フェノール樹脂として、前記の一般式②で 表されるフェノール樹脂を含有するので、硬化物の吸湿 性が小さくなり、密着力が向上し、耐湿性、耐半田クラ ック性に優れる。

【0020】本発明の請求項3に係る半導体装置、請求項1又は請求項2記載の封止用エポキシ樹脂組成物の硬化物により封止してなるので、耐湿性に優れ、緑色であり、ポストキュア一後の変色を低減できる。 【0021】

【実施例】以下、本発明を実施例によって具体的に説明 する。

【0022】 (実施例1~実施例3及び比較例1~比較 例4)表1に示した配合で下記の原料を使用した。エポ キシ樹脂として、エポキシ当量195のオルソークレゾ ールノボラック型エポキシ樹脂〔住友化学工業社製;品 番ESCN195 XL-3] (表1でエポキシ樹脂A と記した)、ビフェニル型エポキシ樹脂 (油化シェルエ ポキシ社製;品番YX4000日) (表1でエポキシ樹 脂Bと記した)又はジシクロペンタジエン系エポキシ樹 脂「大日本インキ化学工業社製:品番EXA7200 H) を用いた。硬化剤として、フェノールノボラック樹 脂「荒川化学社製:商品名タマノール752) (表1で 硬化剤Dと記した)又は軟化点80℃の前記の一般式③ で表されるフェノール樹脂 (表1で硬化剤Eと記した) を用いた。充填剤として、 y - グリシドキシプロピルト リメトキシシラン (カップリング剤) でカップリング処 理した溶融シリカを用いた。硬化促進剤として2フェニ ルイミダゾール、雑燃剤としてエポキシ当量400のプ ロム化エポキシ樹脂及び三酸化アンチモン、離型剤とし てカルナバワックスを用いた。顔料としてカーボンブラ ック、塩素化銅フタロシアニングリーン〔ゼネカ社製; 商品名パイナモングリーン〕を用いた。

【0023】 前記の各実施例及び比較例において、前記 配合成分を表1で示した配合の割合で用いてミキサーで 3分間均一に混合分散した後、ロール温度100~12 0℃のミキシングロールで加熱、溶融、混練した。この 混練物を、冷却し、粉砕して各封止用エポキシ樹脂組成 物を得た。

ル骨格を有するため、ピフェニル型エポキシ樹脂を用いることにより、得られる封止品が低弾性率であって、高 スファー成形機を用いて金型温度 175℃、成形圧力5 い熱時強度を持ち、密着力に優れたものになり、ジシク 50 0kg/cm<sup>\*</sup>で半導体素子を封止成形して60QFP TEG (外形:  $15 \, \text{mm} \times 19 \, \text{mm} \times t2$ .  $4 \, \text{mm}$ ) を得た。また、同じ成形条件で $\phi 50 \, \text{mm} \times t3 \, \text{mm}$ の 吸湿率測定用円板及びポストキュア一後の変色測定用円 板を得た。また、密着性測定用の $\phi 11$ .  $3 \, \text{mm} \times t1 \, \text{mm}$ のブリン型の成形品を得た。

【0025】吸湿率、接着強度、耐吸湿半田クラック、

\* 耐湿信頼性及びポストキュアー後の変色を測定した結果、表1に示したように実施例1~実施例3は、比較例1~比較例4より優れていることが確認できた【0026】

【0026】 【表1】

			安装例1	実施例2	実施例3	比較例 1	比較例 2	比較例3	比較例 4
		Cポキシ樹脂 A	-	-	-	119	119	119	119
R		Cボキシ樹脂B	119	-	45	-	-	-	-
	エポキシ樹脂で			130	45	-	-	-	Н
	硬化射D		67	56	-	67	67	87	67
		硬化射 E		-	96		-	-	-
	プロム化エポキシ樹脂		14	14	14	14	14	14	14
a	三酸化アンチモン		14.5	14.5	14.5	14.5	14.5	14.5	14.5
	硬化促進劑		2	2	2	2	2	2	2
_		#型剂	3	3	3	3	3	3	3
	×	イナモングリーン	5	5	5	0.5	10	5	-
#	カーポンプラック		0.5	0.5	0.5		2		5
%	نا	カップリング剣	6	5	5	5	5	5	- 5
_		溶験シリカ	770	770	7.70	775	763.5	770.5	770.5
		合計	1000	1000	1000	1000	1000	1000	1000
	0	温率 [重量%]	0.18	0.21	0.14	0.30	0.30	0.30	0.30
拝	ほ選半田クラック 10個中の不良機			3		10	10	10	10
	附信	200時間後の 10個中の不良数		7		10	10	10	10
	進性	500時間後の 10個中の不良数	5	10		_	_	_	-
	密滑性[kgf/cm²]		45	35	47	12	12	12	12
		トキュアー後の変色	角好	章杆	島軒	不可	B 147	不可	由好

【0027】表1において記載した測定値は、次の方法によった。

# (1)吸湿率

JIS K 6911準じて作製した直径50mm、厚み3mmの円板を温度85℃、相対湿度85%の雰囲気に放置し、72時間後の重量変化を測定した。

#### (2)耐吸湿半田クラック

60QFP TECを温度85℃、相対湿度85%の雰囲気に放置し、72時間吸湿後、250℃の半田に10 砂間浸漬し、実体脈微鏡でクラックの有無を観察し、6 QFP10個中でクラックが発生したパッケージの個数を求めた。

#### (3) 耐湿信頼性

600FP TECを温度 5 °C、相対温度 8 5 %の雰囲気に放置し、7 2 時間吸温後、2 5 0 °Cの半田に 10 秒間浸漬し、PCT (プレッシャークッカーテスト) 1 3 3 °C、相対温度 10 0 %の雰囲気に放置し、2 0 0 時 40 間及び 5 0 0 時間後のアルミ回路のオープン不良(回路 断線)の有無を観察し、6 0 0 FP 1 0 個中でオープン不良が発生したパッケージの個数を求めた。

# (4)密着性

ブリン型のキャピティにニッケル板で蓋をし、得られた 封止用エポキン樹脂組成物を170°でに加熱した金型内 にトランスファー注入して硬化させ、トランスファー成 形で成形品を得て、ニッケル板と成形品との密着力をブ ッシュブルゲージで測定した。

#### (5) ポストキュアー後の変色

175℃、6時間のポストキュアー前と後との色差を色差計で測定し、色差ΔEが2以下の場合を良好、2を越える場合を不可とした。

#### [0028]

【発明の効果】本発明の請求項1に係る封止用エポキシ 棚別組成物によると、ピフェニル型エポキン樹脂的側面 なピフェニル骨格を有するため、ピフェニル型エポキシ 樹脂を用いることにより、得られる封止品が低弾性率で あって、高い熱時強度を持ち、密着力に優れたものにな・ り、ジシクロペンタジェン系エポキシ樹脂を用いること により、硬化物の吸湿性が小さくなる。かつ、前記餌料 として、塩素化銅フタロシアニングリーンを対止用エポ キシ樹脂組が全量に対して0、05~1 電量及及びカ ーポンブラックを封止用エポキシ樹脂組成物全量に対し て0・01~0、20重像と含有するので、ポストキュア 一後の変色を低減した緑色の対止用エポキン樹脂組成物 が得られる。すなわち、耐湿性、耐吸湿半田クラック 性、密着性に優れ、ポストキュア一後の変色を低減した 緑色の成形品が得られる。

【0029】本発明の請求項2に係る封止用エポキシ樹 脂組成物によると、吸湿性が小さくなり、密着力が向上 し、耐湿性、耐半田クラック性に優れた成形品が得られる。

【0030】本発明の請求項3に係る半導体装置による と、請求項1又は請求項2記載の封止用エポキシ樹脂組 成物の硬化物により封止してなるので、耐湿性に優か。 終色であり、ポストキュアー後の変色を低減できる。 フロントページの続き

(72)発明者 池田 博則 大阪府門真市大字門真1048番地松下電工株 式会社内 (72)発明者 池田 幸司 大阪府門真市大字門真1048番地松下電工株 式会社内

(72)発明者 山中 浩史 大阪府門真市大字門真1048番地松下電工株 式会社内

# PATENT ABSTRACTS OF JAPAN

(11)Publication number:

09-008178

(43) Date of publication of application: 10.01.1997

(51)Int.Cl.

H01L 23/29

H01L 23/31

C08G 59/18

C08G 59/26

C08L 63/00

(21)Application number: 07-147661

(71)Applicant: MATSUSHITA ELECTRIC WORKS LTD

(22)Date of filing:

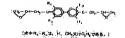
14.06.1995

(72)Inventor: HARA RYUZO

MIVATANI VOSHIHIRO ICHIKAWA TAKAYUKI IKEDA HIRONORI IKEDA KOJI

YAMANAKA HIROSHI

# (54) EPOXY RESIN COMPOSITION FOR SEALING AND SEMICONDUCTOR DEVICE ÙSÍNG IT



(57) Abstract:

PURPOSE: To obtain an epoxy resin composition by which a moisture proof property and an absorbent solder crackresistant property are enhanced by a method wherein a specific biphenyl-type epoxy resin and a dicyclopentadienebased epoxy resin are contained as epoxy resins and chlorinated copper phthalocyanine green and carbon black in respectively specific amounts are contained as pigments. CONSTITUTION: A biphenyl-type epoxy resin expressed by Formula I and/or a dicyclopentadiene-based epoxy resin expressed by Formula II are contained as epoxy resins. Since the biphenyl-type epoxy resin comprises a rigid framework, a sealed product which is obtained has a low modulus of elasticity, its strength is high at a high temperature, and its close contact property is excellent. In addition, the absorbency of a hardened substance is small because the dicyclopentadiene-based epoxy resin is contained. In addition, 0.05 to 1wt.% of chlorinated copper



phthalocyanine green with reference to the total amount of a resin component and 0.01 to 0.2wt.% of carbon black

with reference to the total amount are contained as pigments. As a result, it is possible to reduce a change of color after a posturing operation.

[Date of request for examination]

Date of sending the examiner's decision of rejection1

[Kind of final disposal of application other than the examiner's decision of rejection or application converted registration

[Date of final disposal for application]

[Patent number]

[Date of registration]

[Number of appeal against examiner's decision of rejection

Date of requesting appeal against examiner's

decision of rejection]

[Date of extinction of right]

# CLAIMS

# [Claim(s)]

[Claim 1] The phenol resin which has two or more phenolic hydroxyl groups in 1 molecule as a curing agent in the epoxy resin which has two or more epoxy groups in 1 molecule. In the epoxy resin constituent for the closures which comes to add an inorganic bulking agent, a hardening accelerator, and a pigment The dicyclopentadiene system epoxy resin expressed with the biphenyl mold epoxy resin and/or following general formula \*\* which are expressed with following general formula \*\* as said epoxy resin is contained. As said pigment The epoxy resin constituent for the closures characterized by containing 0.05 - 1 % of the weight, and carbon black for chlorination copper Phthalocyanine Green 0.01 to 0.2% of the weight to the epoxy resin constituent whole quantity for the closures to the epoxy resin constituent whole quantity for the closures. [Formula 1]

(式中R1~R4は、H、CH4又はC4H2である。)

(式中mは繰り返し単位を示す0~5の数)

[Claim 2] The epoxy resin constituent for the closures according to claim 1 characterized by containing the phenol resin expressed with following general formula \*\* as said phenol resin. [Formula 31

[Claim 3] The semiconductor device characterized by coming to close with the hardened material of the epoxy resin constituent for the closures according to claim 1 or 2.

# DETAILED DESCRIPTION

# [Detailed Description of the Invention]

[0001]

[Industrial Application] This invention relates to the epoxy resin constituent for the closures for closing an electrical part, electronic parts, a semiconductor device, etc., and the semiconductor device using it.

[0002]

[Description of the Prior Art] Although the hermetic-sealing method using the resin seal approach by the epoxy resin, silicon resin, etc., glass, a metal, the ceramics, etc. has been conventionally adopted as the closure approaches, such as electrical and electric equipment and electronic parts, such as diode, a transistor, and an integrated circuit, and a semiconductor device In the closure method using the epoxy resin which has a merit in respect of mass production method or cost with improvement in dependability in recent years Most generally the molding material which consists of a constituent which uses cresol novolak mold resin as a resinous principle, and uses phenol novolak mold resin as a curing agent component is used.

[0003] The thinning of mold resin is progressing and it is impossible however, to not necessarily correspond to satisfaction with an old epoxy resin constituent with the densification of electronic parts, such as IC, LSI, and VLSI, or a semiconductor device, and high integration. For example, in the device for surface mounts, since the device itself is rapidly put [ immerse / in solder / directly ] to the bottom of an elevated-temperature cruel environment at the time of mounting, it has been the situation under which generating of a package crack is not avoided. That is, a crack arises in the package of a semiconductor device, without the moisture which absorbed moisture during the storage after shaping carrying out evaporation expansion rapidly [ in case it is exposed to an elevated temperature ], and closure resin being able to finishing bearing this. Moreover, although there are some for which green is used in some of diodes and transistors, generally the epoxy resin constituent for the closures is black. There was a trouble that want to use it as an epoxy resin constituent for the closures since an image is good and conspicuous, but moisture resistance of green would be bad and would discolor it after a post cure if the usual green pigment is used. [0004]

[Problem(s) to be Solved by the Invention] The place which this invention was made in view of the aforementioned situation, and is made into the purpose is excellent in moisture resistance, moisture absorption-proof solder crack nature, and adhesion, and is to offer the semiconductor device using the green epoxy resin constituent for the closures and green it which reduced the discoloration after a post cure.

T00051

[Means for Solving the Problem] The epoxy resin constituent for the closures concerning claim 1 of this invention The phenol resin which has two or more phenolic hydroxyl groups in 1 molecule as a

curing agent in the epoxy resin which has two or more epoxy groups in 1 molecule, In the epoxy resin constituent for the closures which comes to add an inorganic bulking agent, a hardening accelerator, and a pigment The dicyclopentadiene system epoxy resin expressed with the biphenyl mold epoxy resin and/or following general formula \*\* which are expressed with following general formula \*\* as said epoxy resin is contained. As said pigment It is characterized by containing 0.05 - 1 % of the weight, and carbon black for chlorination copper Phthalocyanine Green 0.01 to 0.2% of the weight to the epoxy resin constituent whole quantity for the closures to the epoxy resin constituent whole quantity for the closures.

[0006]

(式中R1~R2は、H、CH3又はC3H2である。)

(式中mは繰り返し単位を示す0~5の数)

[0008] The epoxy resin constituent for the closures concerning claim 2 of this invention is characterized by containing the phenol resin expressed with following general formula \*\* as said phenol resin.

[0009] [Formula 6]

$$\begin{array}{c|c} & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & &$$

[0010] It is characterized by coming to close with the hardened material of the semiconductor device concerning claim 3 of this invention, and the epoxy resin constituent for the closures according to claim 1 or 2.

[0011] Hereafter, this invention is explained in full detail. The epoxy resin used for this invention is an epoxy resin which has at least two epoxy groups in 1 molecule, and it is indispensable to contain the dicyclopentadiene system epoxy resin expressed with the biphenyl mold epoxy resin and/or above general formula \*\* which are expressed with above general formula \*\* as this epoxy resin. That is, since a biphenyl mold epoxy resin has an upright biphenyl frame, the closure article obtained is a low modulus of elasticity, it has reinforcement at the time of high heat, and becomes the thing excellent in the adhesion force, and the hygroscopicity of a hardened material becomes small by using a dicyclopentadiene system epoxy resin. That is, by using said epoxy resin, moisture absorption can be lowered, the adhesion force with a leadframe can be raised, and it becomes the

epoxy resin constituent for the closures with which the hardened material excellent in moisture resistance and solder-proof crack nature is obtained.

[0012] If it is the curing agent which has at least two phenolic hydroxyl groups in 1 molecule as a curing agent used for this invention, any curing agent an be used, for example, there are phenol novolak resin, cresol novolak resin, etc.

[0013] In this invention, as an inorganic filler, fused silica, a crystal silica, an alumina, silicon nitride, etc. can be used, and diazabicycloundecen, triphenyl phosphine, tetra-phenyl phosphonium, tetraphenyl borate, an imidazole, tertiary amine, etc. can be used as a hardening accelerator. Moreover, release agents, such as stearic acid, can be used for flame retarders, such as coupling agents, such as an epoxy silane, a bromine-ized epoxy resin, and an antimony trioxide, a silicone flexibilizer, and a carmauba wax list if needed.

[0014] It is indispensable to contain chlorination copper Phthalocyanine Green 0.05 to 1% of the weight to the epoxy resin constituent whole quantity for the closures as a pigment in the epoxy resin constituent for the closures concerning this invention. That is, when green becomes thin too much, it becomes impossible to recognize it as green to less than 0.05% of the weight of a case and the content of chlorination copper Phthalocyanine Green exceeds 1 % of the weight to the epoxy resin constituent whole quantity for the closures, moisture resistance falls. It is necessary to contain carbon black 0.01 to 0.2% of the weight to the epoxy resin constituent whole quantity for the closures with chlorination copper Phthalocyanine Green. That is, when discoloration occurs after a post cure in less than 0.01% of the weight of a case to the epoxy resin constituent whole quantity for the closures and the content of carbon black exceeds 0.2 % of the weight, it will become black, without the ability recognizing it as green.

[0015] As for the epoxy resin constituent for the closures concerning this invention, it is desirable to contain the phenol resin expressed with above general formula \*\* as phenol resin which is a curing agent. That is, by using the phenol resin expressed with above general formula \*\*, the hygroscopicity of a hardened material becomes small, the adhesion force improves, and the epoxy resin constituent for the closures which is excellent in moisture resistance and solder-proof crack nature is obtained.

[0016] Thus, it adds a curing agent, an inorganic bulking agent, a hardening accelerator, and a pigment and grinds [mix, knead and] to an epoxy resin, it corns if needed further, and the epoxy resin constituent for the closures is obtained. Furthermore, this epoxy resin constituent for the closures is used, a semiconductor device is closed by transfer molding etc., and a semiconductor device is obtained.

[0017] As mentioned above, according to this invention, it excels in moisture resistance, moisture absorption-proof solder crack nature, and adhesion, and the semiconductor device using the green epoxy resin constituent for the closures and green it which reduced the discoloration after a post cure is obtained.

f00181

[Function] The epoxy resin constituent for the closures concerning claim 1 of this invention Since the dicyclopentadiene system epoxy resin expressed with the biphenyl mold epoxy resin and/or above general formula \*\* which are expressed with above general formula \*\* as an epoxy resin is contained Since it has a biphenyl frame with an upright biphenyl mold epoxy resin, by using a biphenyl mold epoxy resin The closure article obtained is a low modulus of elasticity, it has reinforcement at the time of high heat, and becomes the thing excellent in the adhesion force, and the hygroscopicity of a hardened material becomes small by using a dicyclopentadiene system epoxy resin. And since 0.05 - 1 % of the weight and carbon black are contained for chlorination copper Phthalocyanine Green 0.01 to 0.2% of the weight to the epoxy resin constituent whole quantity for the closures as said pigment to the epoxy resin constituent whole quantity for the closures, the green epoxy resin constituent for the closures which reduced the discoloration after a post cure is obtained.

[0019] Since the epoxy resin constituent for the closures concerning claim 2 of this invention contains the phenol resin expressed with above general formula \*\* as phenol resin, the

hygroscopicity of a hardened material becomes small, and its adhesion force improves, and it is excellent in moisture resistance and solder-proof crack nature.

[0020] Since it comes to close with the hardened material of the semiconductor device concerning claim 3 of this invention, and the epoxy resin constituent for the closures according to claim 1 or 2, it excels in moisture resistance, and it is green and the discoloration after a post cure can be reduced. [0021]

[Example] Hereafter, an example explains this invention concretely.

[0022] (An example 1 - an example 3, and the example 1 of a comparison - the example 4 of a comparison) The raw material following by the combination shown in Table 1 was used, as an epoxy resin -- the polyglycidyl ether of o-cresol-form aldevde novolac of weight per epoxy equivalent 195 -- [ --; lot number ESCNby Sumitomo Chemical Co., Ltd. 195 XL-3] (it was described as epoxy resin A in Table 1), and a biphenyl mold epoxy resin -- [ --; lot number YXby oil-ized shell epoxy company4000H] (it was described as epoxy resin B in Table 1), or a dicyclopentadiene --; lot number EXAby system epoxy resin [Dainippon Ink & Chemicals, Inc.7200H] was used. As a curing agent, the phenol resin (it was described as the curing agent E in Table 1) expressed with aforementioned general formula [ of phenol novolak resin [: trade name TAMANORU 752 by the Arakawa chemistry company] (it was described as the curing agent D in Table 1) or 80 degrees C of softening temperatures 1 \*\* was used. As a bulking agent, the fused silica which carried out coupling processing by gamma-glycidoxypropyltrimetoxysilane (coupling agent) was used. 2-phenylimidazole was used as a hardening accelerator and carnauba wax was used as a flame retarder as the bromine-ized epoxy resin of weight per epoxy equivalent 400 and an antimony trioxide, and a release agent. Carbon black and chlorination copper Phthalocyanine Green I; trade name BAINAMON Green by Zeneka Co.1 were used as a pigment.

[0023] a roll mill with a roll temperature [ after the combination shown in Table 1 coming out comparatively, using said combination component in each aforementioned example and the aforementioned example of a comparison and carrying out mixed distribution for 3 minutes by the mixer at homogeneity ] of 100-120 degrees C -- heating and melting -- it kneaded. This kneading object was cooled and ground and each epoxy resin constituent for the closures was obtained. [0024] A transfer-molding machine is used for each epoxy resin constituent obtained above, and they are the die temperature of 175 degrees C, and the compacting pressure of 50kg/cm2. Closure shaping of the semiconductor device is carried out, and it is 60QFP. TEG (appearance: 15mmx19 mmxt2.4mm) was obtained. Moreover, the disk for moisture absorption measurement of phi50 mmxt3mm and the disk for discoloration measurement after a post cure were obtained by the same process condition. Moreover, the mold goods of the pudding mold of phi11.3 mmxt10mm for adhesion measurement were obtained.

[0025] As a result of measuring the discoloration after moisture absorption, bond strength, a moisture absorption-proof solder crack, humidity-tolerant reliability, and a post cure, it is [0026] which has checked that the example 1 - the example 3 were superior to the example 1 of a comparison - the example 4 of a comparison as shown in Table 1.

[Table 1]

			実施例1	実施例2	実施例3	比較例1	比較例 2	比較例3	比較例 4
Т		エポキシ樹脂A			-	119	119	119	119
配		エポキシ樹脂B	119	_	45		_	_	_
[		エポキシ樹脂で		130	45	-	_	_	_
		硬化剤D	67	56	-	67	87	67	67
L		硬化剤 E			96	1	-	-	
L	ブロ	ム化エポキシ樹脂	14	14	14	14	14	14	14
슴	=	酸化アンチモン	14.5	14.5	14.5	14.5	14.5	14.5	14.5
- 1		硬化促進剤	2	2	2	2	2	2	2
-1	_	能型剂	3	3	3	3	3	3	3
=	14.	イナモングリーン	5	5	5	0.5	10	5	-
=		ーポンプラック	0.5	0.5	0.5	-	2		5
%[		カップリング剤	5	5	5	5	5	5	5
띡		溶融シリカ	770	770	7,70	775	763.5	770.5	770.5
_[	合計		1000	1000	1000	1000	1000	1000	1000
╗	9	[聖皇 [重量%]	0.18	0.21	0.14	0.30	0.30	0.30	0.30
۳		選半田クラック 10個中の不良数	٥	3	0	10	10	10	10
ſ	推論	200時間後の 10個中の不良数	٥	7	0	10	10	10	10
1	湿性	500時間後の 10個中の不良数	5	10	0	_	_	-	_
循	- 6	Brite [kgf/cm*]	45	35	47	12	12	12	12
_[	ボス	トキュアー後の変色	良好	良好	良奸	不可	泉好	不可	良好

[0027] The measured value indicated in Table 1 was based on the following approach.

- (1) Moisture absorption JIS K It applied correspondingly 6911, the produced disk with a diameter [of 50mm] and a thickness of 3mm was left in the ambient atmosphere of the temperature of 85 degrees C, and 85% of relative humidity, and weight change 72 hours after was measured.
- (2) Moisture absorption-proof solder crack 60QFP It asked for the number of the package which left TEG in the ambient atmosphere of the temperature of 85 degrees C, and 85% of relative humidity, was immersed in 250-degree C solder for 10 seconds after 72-hour moisture absorption, and observed the existence of a crack with the stereoscopic microscope, and the crack generated in ten 60QFP.
- (3) Humidity-tolerant reliability 60QFP TEG was left in the ambient atmosphere of the temperature of 85 degrees C, and 85% of relative humidity, and it was immersed in 250-degree C solder for 10 seconds after 72-hour moisture absorption, and was left in PCT(pressure cooker test) 133degree C and the ambient atmosphere of 100% of relative humidity, the existence of poor opening (circuit open circuit) of the aluminum circuit of 200 hours and 500 hours after was observed, and it asked for the number of the package which poor opening generated in ten 60QFP.
- (4) The cavity of an adhesion pudding mold was covered with the nickel plate, in the metal mold heated at 170 degrees C, transfer impregnation is carried out, the obtained epoxy resin constituent for the closures was stiffened, mold goods were obtained by transfer molding, and the adhesion force of a nickel plate and mold goods was measured with the push pull gage.
- (5) 175 degrees C of discoloration after a post cure and the color difference of the post cure front of 6 hours and the back were measured with the color difference meter, and the case where fitness and 2 were exceeded for the case where color difference deltaE is two or less was made improper. [0028]

[Effect of the Invention] Since it has a biphenyl frame with an upright biphenyl mold epoxy resin according to the epoxy resin constituent for the closures concerning claim 1 of this invention, by using a biphenyl mold epoxy resin, the closure article obtained is a low modulus of elasticity, it has reinforcement at the time of high heat, and becomes the thing excellent in the adhesion force, and the hygroscopicity of a hardened material becomes small by using a dicyclopentadiene system epoxy resin. And since 0.05 - 1 % of the weight and carbon black are contained for chlorination copper Phthalocyanine Green 0.01 to 0.2% of the weight to the epoxy resin constituent whole quantity for the closures as said pigment to the epoxy resin constituent whole quantity for the closures, the green epoxy resin constituent for the closures which reduced the discoloration after a post cure is obtained. That is, it excels in moisture resistance, moisture absorption-proof solder

crack nature, and adhesion, and the green mold goods which reduced the discoloration after a post cure are obtained.

[0029] According to the epoxy resin constituent for the closures concerning claim 2 of this invention, hygroscopicity becomes small, the adhesion force improves and mold goods excellent in moisture resistance and solder-proof crack nature are obtained.

[0030] Since it comes to close with the hardened material of the epoxy resin constituent for the closures according to claim 1 or 2 according to the semiconductor device concerning claim 3 of this invention, it excels in moisture resistance, and it is green and the discoloration after a post cure can be reduced.

# TECHNICAL FIELD

[Industrial Application] This invention relates to the epoxy resin constituent for the closures for closing an electrical part, electronic parts, a semiconductor device, etc., and the semiconductor device using it.

# PRIOR ART

[Description of the Prior Art] Although the hermetic-sealing method using the resin seal approach by the epoxy resin, silicon resin, etc., glass, a metal, the ceramics, etc. has been conventionally adopted as the closure approaches, such as electrical and electric equipment and electronic parts, such as diode, a transistor, and an integrated circuit, and a semiconductor device In the closure method using the epoxy resin which has a merit in respect of mass production method or cost with improvement in dependability in recent years Most generally the molding material which consists of a constituent which uses cresol novolak mold resin as a resinous principle, and uses phenol novolak mold resin as a curing agent component is used.

[0003] The thinning of mold resin is progressing and it is impossible however, to not necessarily correspond to satisfaction with an old epoxy resin constituent with the densification of electronic parts, such as IC, LSI, and VLSI, or a semiconductor device, and high integration. For example, in the device for surface mounts, since the device itself is rapidly put [ immerse / in solder / directly ] to the bottom of an elevated-temperature cruel environment at the time of mounting, it has been the situation under which generating of a package crack is not avoided. That is, a crack arises in the package of a semiconductor device, without the moisture which absorbed moisture during the storage after shaping carrying out evaporation expansion rapidly [ in case it is exposed to an elevated temperature ], and closure resin being able to finishing bearing this. Moreover, although there are some for which green is used in some of diodes and transistors, generally the epoxy resin constituent for the closures is black. There was a trouble that want to use it as an epoxy resin constituent for the closures since an image is good and conspicuous, but moisture resistance of green would be bad and would discolor it after a post cure if the usual green pigment is used.

# EFFECT OF THE INVENTION

[Effect of the Invention] Since it has a biphenyl frame with an upright biphenyl mold epoxy resin according to the epoxy resin constituent for the closures concerning claim 1 of this invention, by using a biphenyl mold epoxy resin, the closure article obtained is a low modulus of elasticity, it has reinforcement at the time of high heat, and becomes the thing excellent in the adhesion force, and the hygroscopicity of a hardened material becomes small by using a dicyclopentadiene system epoxy resin. And since 0.05 - 1 % of the weight and carbon black are contained for chlorination copper Phthalocyanine Green 0.01 to 0.2% of the weight to the epoxy resin constituent whole

quantity for the closures as said pigment to the epoxy resin constituent whole quantity for the closures, the green epoxy resin constituent for the closures which reduced the discoloration after a post cure is obtained. That is, it excels in moisture resistance, moisture absorption-proof solder crack nature, and adhesion, and the green mold goods which reduced the discoloration after a post cure are obtained

[0029] According to the epoxy resin constituent for the closures concerning claim 2 of this invention, hygroscopicity becomes small, the adhesion force improves and mold goods excellent in moisture resistance and solder-proof crack nature are obtained.

[0030] Since it comes to close with the hardened material of the epoxy resin constituent for the closures according to claim 1 or 2 according to the semiconductor device concerning claim 3 of this invention, it excels in moisture resistance, and it is green and the discoloration after a post cure can be reduced.

# TECHNICAL PROBLEM

[Problem(s) to be Solved by the Invention] The place which this invention was made in view of the aforementioned situation, and is made into the purpose is excellent in moisture resistance, moisture absorption-proof solder crack nature, and adhesion, and is to offer the semiconductor device using the green epoxy resin constituent for the closures and green it which reduced the discoloration after a post cure.

# MEANS

[Means for Solving the Problem] The epoxy resin constituent for the closures concerning claim 1 of this invention The phenol resin which has two or more phenolic hydroxyl groups in 1 molecule as a curing agent in the epoxy resin which has two or more epoxy groups in 1 molecule. In the epoxy resin constituent for the closures which comes to add an inorganic bulking agent, a hardening accelerator, and a pigment The dicyclopentadiene system epoxy resin expressed with the biphenyl mold epoxy resin and/or following general formula \*\* which are expressed with following general formula \*\* as said epoxy resin is contained. As said pigment It is characterized by containing 0.05 -1 % of the weight, and carbon black for chlorination copper Phthalocyanine Green 0.01 to 0.2% of the weight to the epoxy resin constituent whole quantity for the closures to the epoxy resin constituent whole quantity for the closures.

(式中R,-R,は、H、CH,又はC,H,である。)

(式中mは繰り返し単位を示す0~5の数)

[0008] The epoxy resin constituent for the closures concerning claim 2 of this invention is characterized by containing the phenol resin expressed with following general formula \*\* as said phenol resin.

[0009] [Formula 6]

[0010] It is characterized by coming to close with the hardened material of the semiconductor device concerning claim 3 of this invention, and the epoxy resin constituent for the closures according to claim 1 or 2.

[0011] Hereafter, this invention is explained in full detail. The epoxy resin used for this invention is an epoxy resin which has at least two epoxy groups in 1 molecule, and it is indispensable to contain the dicyclopentadiene system epoxy resin expressed with the biphenyl mold epoxy resin and/or above general formula \*\* which are expressed with above general formula \*\* as this epoxy resin. That is, since a biphenyl mold epoxy resin has an upright biphenyl frame, the closure article obtained is a low modulus of elasticity, it has reinforcement at the time of high heat, and becomes the thing excellent in the adhesion force, and the hygroscopicity of a hardened material becomes small by using a dicyclopentadiene system epoxy resin. That is, by using said epoxy resin, moisture absorption can be lowered, the adhesion force with a leadframe can be raised, and it becomes the epoxy resin constituent for the closures with which the hardened material excellent in moisture resistance and solder-proof crack nature is obtained.

[0012] If it is the curing agent which has at least two phenolic hydroxyl groups in 1 molecule as a curing agent used for this invention, any curing agent can be used, for example, there are phenol novolak resin, cresol novolak resin, polyfunctional phenol resin, etc.

[0013] In this invention, as an inorganic filler, fused silica, a crystal silica, an alumina, silicon nitride, etc. can be used, and diazabicycloundecen, triphenyl phosphine, tetra-phenyl phosphonium, tetraphenyl borate, an imidazole, tertiary amine, etc. can be used as a hardening accelerator. Moreover, release agents, such as stearic acid, can be used for flame retarders, such as coupling agents, such as an epoxy silane, a bromine-ized epoxy resin, and an antimony trioxide, a silicone flexibilizer, and a carnauba wax list if needed.

[0014] It is indispensable to contain chlorination copper Phthalocyanine Green 0.05 to 1% of the weight to the epoxy resin constituent whole quantity for the closures as a pigment in the epoxy resin constituent for the closures concerning this invention. That is, when green becomes thin foo much, it becomes impossible to recognize it as green to less than 0.05% of the weight of a case and the content of chlorination copper Phthalocyanine Green exceeds 1 % of the weight to the epoxy resin constituent whole quantity for the closures, moisture resistance falls. It is necessary to contain carbon black 0.01 to 0.2% of the weight to the epoxy resin constituent whole quantity for the closures with chlorination copper Phthalocyanine Green. That is, when discoloration occurs after a post cure in less than 0.01% of the weight of a case to the epoxy resin constituent whole quantity for the closures and the content of carbon black exceeds 0.2 % of the weight, it will become black, without the ability recognizing it as green.

[0015] As for the epoxy resin constituent for the closures concerning this invention, it is desirable to contain the phenol resin expressed with above general formula \*\* as phenol resin which is a curing agent. That is, by using the phenol resin expressed with above general formula \*\*, the

hygroscopicity of a hardened material becomes small, the adhesion force improves, and the epoxy resin constituent for the closures which is excellent in moisture resistance and solder-proof crack nature is obtained.

[0016] Thus, it adds a curing agent, an inorganic bulking agent, a hardening accelerator, and a pigment and grinds [mix, knead and] to an epoxy resin, it corns if needed further, and the epoxy resin constituent for the closures is obtained. Furthermore, this epoxy resin constituent for the closures is used, a semiconductor device is closed by transfer molding etc., and a semiconductor device is obtained.

[0017] As mentioned above, according to this invention, it excels in moisture resistance, moisture absorption-proof solder crack nature, and adhesion, and the semiconductor device using the green epoxy resin constituent for the closures and green it which reduced the discoloration after a post cure is obtained.

# OPERATION

[Function] The epoxy resin constituent for the closures concerning claim 1 of this invention Since the dicyclopentadiene system epoxy resin expressed with the biphenyl mold epoxy resin and/or above general formula \*\* which are expressed with above general formula \*\* as an epoxy resin is contained Since it has a biphenyl frame with an upright biphenyl mold epoxy resin, by using a biphenyl mold epoxy resin. The closure article obtained is a low modulus of elasticity, it has reinforcement at the time of high heat, and becomes the thing excellent in the adhesion force, and the hygroscopicity of a hardened material becomes small by using a dicyclopentadiene system epoxy resin. And since 0.05 - 1 % of the weight and carbon black are contained for chlorination copper Phthalocyanine Green 0.01 to 0.2% of the weight to the epoxy resin constituent whole quantity for the closures as said pigment to the epoxy resin constituent whole quantity for the closures, the green epoxy resin constituent for the closures which reduced the discoloration after a post cure is obtained.

[0019] Since the epoxy resin constituent for the closures concerning claim 2 of this invention contains the phenol resin expressed with above general formula \*\* as phenol resin, the hygroscopicity of a hardened material becomes small, and its adhesion force improves, and it is excellent in moisture resistance and solder-proof crack nature.

[0020] Since it comes to close with the hardened material of the semiconductor device concerning claim 3 of this invention, and the epoxy resin constituent for the closures according to claim 1 or 2, it excels in moisture resistance, and it is green and the discoloration after a post cure can be reduced.

# EXAMPLE

[Example] Hereafter, an example explains this invention concretely.

[0022] (An example 1 - an example 3, and the example 1 of a comparison - the example 4 of a comparison) The raw material following by the combination shown in Table 1 was used. as an epoxy resin -- the polyglycidyl ether of o-cresol-form aldeyde novolac of weight per epoxy equivalent 195 -- [--; lot number ESCNby Sumitomo Chemical Co., Ltd. 195 XL-3] (it was described as epoxy resin A in Table 1), and a biphenyl mold epoxy resin -- [--; lot number YXby oil-ized shell epoxy company4000H] (it was described as epoxy resin B in Table 1), or a dicyclopentadiene --; lot number EXAby system epoxy resin [Dainippon Ink & Chemicals, Inc. 7200H] was used. As a curing agent, the phenol resin (it was described as the curing agent E in Table 1) expressed with aforementioned general formula [ of phenol novolak resin [; trade name TAMANORU 752 by the Arakawa chemistry company] (it was described as the curing agent D in Table 1) or 80 degrees C of softening temperatures ] \*\* was used. As a bulking agent, the fused silica which carried out coupling processing by gamma-glycidoxypropyltrimetoxysilane (coupling

agent) was used. 2-phenylimidazole was used as a hardening accelerator and carnauba wax was used as a flame retarder as the bromine-ized epoxy resin of weight per epoxy equivalent 400 and an antimony trioxide, and a release agent. Carbon black and chlorination copper Phthalocyanine Green [; trade name BAINAMON Green by Zeneka Co.] were used as a pigment.

[0023] a roll mill with a roll temperature [after the combination shown in Table 1 coming out comparatively, using said combination component in each aforementioned example and the aforementioned example of a comparison and carrying out mixed distribution for 3 minutes by the mixer at homogeneity ] of 100-120 degrees C -- heating and melting -- it kneaded. This kneading object was cooled and ground and each epoxy resin constituent for the closures was obtained. [0024] A transfer-molding machine is used for each epoxy resin constituent obtained above, and they are the die temperature of 175 degrees C, and the compacting pressure of 50kg/cm2. Closure shaping of the semiconductor device is carried out, and it is 60QFP. TEG (appearance: 15mmx19 mmxt2.4mm) was obtained. Moreover, the disk for moisture absorption measurement of phi50 mmxt3mm and the disk for discoloration measurement after a post cure were obtained by the same process condition. Moreover, the mold goods of the pudding mold of phi11.3 mmxt10mm for adhesion measurement were obtained.

[0025] As a result of measuring the discoloration after moisture absorption, bond strength, a moisture absorption-proof solder crack, humidity-tolerant reliability, and a post cure, it is [0026] which has checked that the example 1 - the example 3 were superior to the example 1 of a comparison - the example 4 of a comparison as shown in Table 1.

[T	able	e 1]				•			
			実施例1	実脏例2	実施例3	比較例1	比較例 2	比較例3	比較例 4
		エポキシ樹脂A	-		-	119	119	119	119
配		エポキシ樹脂B	119	_	45	-	-	-	~
1		エポキシ樹脂で	-	130	45	_	-	-	-
1	硬化剂D		67	56	-	67	67	67	67
ı		硬化剂E		_	96	_	-	-	-
П	プロ	ム化エポキシ樹脂	14	14	14	14	14	14	14
合	_ =	酸化アンチモン	14.5	14.5	14.5	14.5	14.5	14.5	14.5
ı	L.,	硬化促進剤	2	2	2	2	2	2	2
۱–		#型剂	3	3	3	3	3	3	3
1	``	イナモングリーン	5	5	5	0.5	10	5	_
鼌		カーポンプラック		0.5	0.5	-	2	_	5
%	L_	カップリング剤		5	5	5	5	5	5
l۳		溶験シリカ	770	770	7,70	775	763.5	770.5	770.5
ш		合計		1000	1000	1000	1000	1000	1000
Г		[重量%]	0.18	0.21	0.14	0.30	0.30	0.30	0.30
押		級選半田クラック 10個中の不良数		3	0	10	10	10	10
	耐信額	200時間後の 10個中の不良数	0	7	0	10	10	10	10
	湿性	500時間後の 10個中の不良数	5	10	0	_	_	1	_
衝		唐性[kgf/cm*]	45	35	47	12	12	12	12
	ボス	ポストキュアー後の変色		良好	良好	不可	良好	不可	良好

[0027] The measured value indicated in Table 1 was based on the following approach.

(1) Moisture absorption JIS K It applied correspondingly 6911, the produced disk with a diameter [of 50mm] and a thickness of 3mm was left in the ambient atmosphere of the temperature of 85 degrees C, and 85% of relative humidity, and weight change 72 hours after was measured.

(2) Moisture absorption-proof solder crack 60QFP It asked for the number of the package which left TEG in the ambient atmosphere of the temperature of 85 degrees C, and 85% of relative humidity, was immersed in 250-degree C solder for 10 seconds after 72-hour moisture absorption, and observed the existence of a crack with the stereoscopic microscope, and the crack generated in ten 60QFP.

(3) Humidity-tolerant reliability 60QFP TEG was left in the ambient atmosphere of the temperature of 85 degrees C, and 85% of relative humidity, and it was immersed in 250-degree C solder for 10 seconds after 72-hour moisture absorption, and was left in PCT(pressure cooker test) 133degree C

and the ambient atmosphere of 100% of relative humidity, the existence of poor opening (circuit open circuit) of the aluminum circuit of 200 hours and 500 hours after was observed, and it asked for the number of the package which poor opening generated in ten 600FP.

(4) The cavity of an adhesion pudding mold was covered with the nickel plate, in the metal mold heated at 170 degrees C, transfer impregnation is carried out, the obtained epoxy resin constituent for the closures was stiffened, mold goods were obtained by transfer molding, and the adhesion force of a nickel plate and mold goods was measured with the push pull gage.

(5) 175 degrees C of discoloration after a post cure and the color difference of the post cure front of

6 hours and the back were measured with the color difference meter, and the case where fitness and 2 were exceeded for the case where color difference deltaE is two or less was made improper.